Genestran 75 micrograms/ml solution for injection for cattle, horses and pigs

Authorised

- R-Cloprostenol sodium

Product identification

Medicine name:
Genestran 75 micrograms/ml solution for injection for cattle, horses and pigs
GENESTRAN 75 mikrogramov/ml raztopina za injiciranje za govedo, konje in prašiče

Active substance:
R-Cloprostenol sodium

Target species:
Cattle
Horse
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
R-Cloprostenol sodium

78.88 microgram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Meat and offal. 1 day
  - Milk. 0 day
- **Horse**
  - Meat and offal. 1 day
  - Milk. 0 hour
- **Pig**
  - Meat and offal. 1 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QG02AD90

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Slovenia

Available in:
Slovenia

Package description:
Colourless vial of type I glass containing 20 ml of solution for injection, with chlorobutyl rubber stopper and aluminium cap.Presentation: Cardboard box of 1 vial

of 20 ml
Colourless vial of type I glass containing 50 ml of solution for injection, with chlorobutyl rubber stopper and aluminium cap.Presentation: Cardboard box of 1 vial of 50 ml
Colourless vial of type I glass containing 20 ml of solution for injection, with chlorobutyl rubber stopper and aluminium cap.Presentation: Cardboard box of 5 vials of 20 ml

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:
aniMedica GmbH

Marketing authorisation date:
14/06/2010

Manufacturing sites for batch release:
aniMedica GmbH
Industrial Veterinaria S.A.

Responsible authority:
Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:
MR/V/0153/001

Date of authorisation status change:
14/06/2010

Reference member state:
Ireland

Procedure number:

IE/V/0228/001

Concerned member states:
Austria Belgium Czechia Estonia France Germany Iceland Italy Latvia Lithuania Luxembourg Norway Poland Portugal Romania Slovakia Slovenia Spain United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

English (PDF)
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Package Leaflet

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Labelling

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